In the claims

The following amendments are made with respect to the claims in the International application PCT/GB2003/005286.

This listing of claims will replace all prior versions and listings of claims in this application.

- 1 (Currently amended). A process for increasing the optical purity of a mixture of enantiomers of mefloquine, wherein said process comprises using a substantially single enantiomer of a O,O-di-p-aroyltartaric acid as a resolving agent.
- 2 (Currently amended). [[A]] <u>The</u> process according to claim 1, for preparing a substantially single enantiomer of mefloquine, which proceeds by means of resolution of racemic mefloquine using a substantially single enantiomer of a O,O-di-p-aroyltartaric acid as a resolving agent.
- 3 (Currently amended). [[A]] <u>The process according to claim 1 or claim 2</u>, wherein the resolving agent is O,O-di-*p*-toluoyl-L-tartaric acid.
- 4 (Currently amended). [[A]] <u>The</u> process according to any preceding claim <u>1</u>, wherein the mefloquine is contaminated with *threo*-mefloquine.
- 5 (Currently amended). [[A]] <u>The process according to any preceding claim 1</u>, which is conducted in a solvent selected from <u>the group consisting of esters</u>, ketones and halogenated solvents.
- 6 (Currently amended). [[A]] <u>The process according to any preceding claim 1</u>, wherein the resolving agent is present in a sub-stoichiometric quantity, whereby an enantiomer of *erythro*-mefloquine is preferentially obtained.
- 7 (Currently amended). [[A]] <u>The</u> process according to claim 6, which is conducted in the presence of an additional chiral or achiral acid.

8 (Currently amended). [[A]] <u>The</u> process according to <u>any preceding</u> claim <u>1</u>, which further comprises conversion of the salt obtained by the resolution to the free base form of mefloquine or a pharmaceutically acceptable salt thereof.